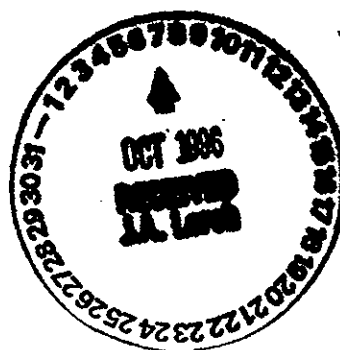


Environmental  
Restoration  
Contractor

**ERC Team**

**Interoffice Memorandum**



0052018

Job No. 22192  
Written Response Required? NO  
Closes CCN: N/A  
OU: 300-FF-2  
TSD: 618 BG / 316-4  
ERA: WELL 699-S6-E4A  
Subject Code: 8660

TO: L C Hulstrom H9-11

DATE: October 7, 1996

COPIES: J A Lerch  
T D Lefrancois  
I D Jacques

FROM: R G McCain  
Env Svcs / OSM Team  
H9-10 / 372-9593

SUBJECT: **FIELD SCREENING RESULTS FOR ORGANIC MATERIAL IN WATER**

46827

On September 19 and 27, 1986, water samples were collected from well 699-S6-E4A, located in the vicinity of the 618 burial ground and 316-4 crib, north of the 300 area. Previous experience with this well had indicated that the water may contain an unknown organic substance.

When the well was opened on September 18, an OVM was used to obtain readings of 6.2-6.6 ppm (isobutylene equivalent) in the vapor space inside the casing. A 5-liter Tedlar bag was collected using the OVM as a sample pump and colorimetric tubes were used to attempt to identify the gas, but results were inconclusive.

On September 19, a preliminary water sample was obtained by bailing the well. Approximately 275 ml of this sample was placed in a one pint (473 ml) wide mouth canning jar and tested for volatile organic compounds using an equilibrium headspace method. In this method, the headspace in the jar is circulated through a photoionization detector (PID). Since a PID is essentially non-destructive, the headspace concentration tends to quickly reach an equilibrium value which can be correlated to the concentration of volatile organic compounds (VOCs) in the water. For the September 19 sample, an OVM was used to measure headspace concentrations. Equilibrium headspace values of 3.9-5.0 ppm (isobutylene equivalent) were observed. This result indicated that the water did contain a VOC detectable with a PID, and plans were made to conduct equilibrium headspace measurements as the well was developed in order to monitor the VOC concentration as a function of time and purge volume.

A downhole pump was installed in well 699-S6-E4A, and samples were collected during well development on September 27, 1996. Screening samples were collected at intervals during well development. Equilibrium headspace measurements are shown on Table 1 (attached)

Both a ThermoEnvironmental OVM and a PhotoVac MicroTip ( $\mu$ TIP) were used to make equilibrium headspace measurements on September 27. The  $\mu$ TIP was added to the test in addition to the OVM because experience has shown that it tends to be more sensitive at low VOC concentrations. This was confirmed by the initial results. Unfortunately, the  $\mu$ TIP drew in water after the third sample and was no longer functional. Both OVMs used for the headspace measurements failed to respond to the VOC present in the water. Hence, little useful data on VOC concentration as a function of development time was obtained.

L C Hulstrom H9-11

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Examination of the water samples did reveal other characteristics of interest. Sample B0J281 was observed to have a pH of 9-10, using pH paper, and the conductivity was 412  $\mu\text{S}/\text{cm}$  at 19.5°C. A small amount of Sudan IV was added to this sample and shaken. After centrifuging for 5 min at 2000 rpm, there was no evidence of any nonaqueous phase. Sudan IV is insoluble in water and soluble in most organic compounds. The presence of an organic nonaqueous phase liquid in the water sample would be expected to produce a bright red layer after centrifuging.

Later samples appeared to contain small amounts of a whitish material which tends to settle out. These samples also had an odor which resembled diesel fuel. The whitish material did not show an affinity for Sudan IV. pH of the later samples was also in the range of 9-10.

R G McCain

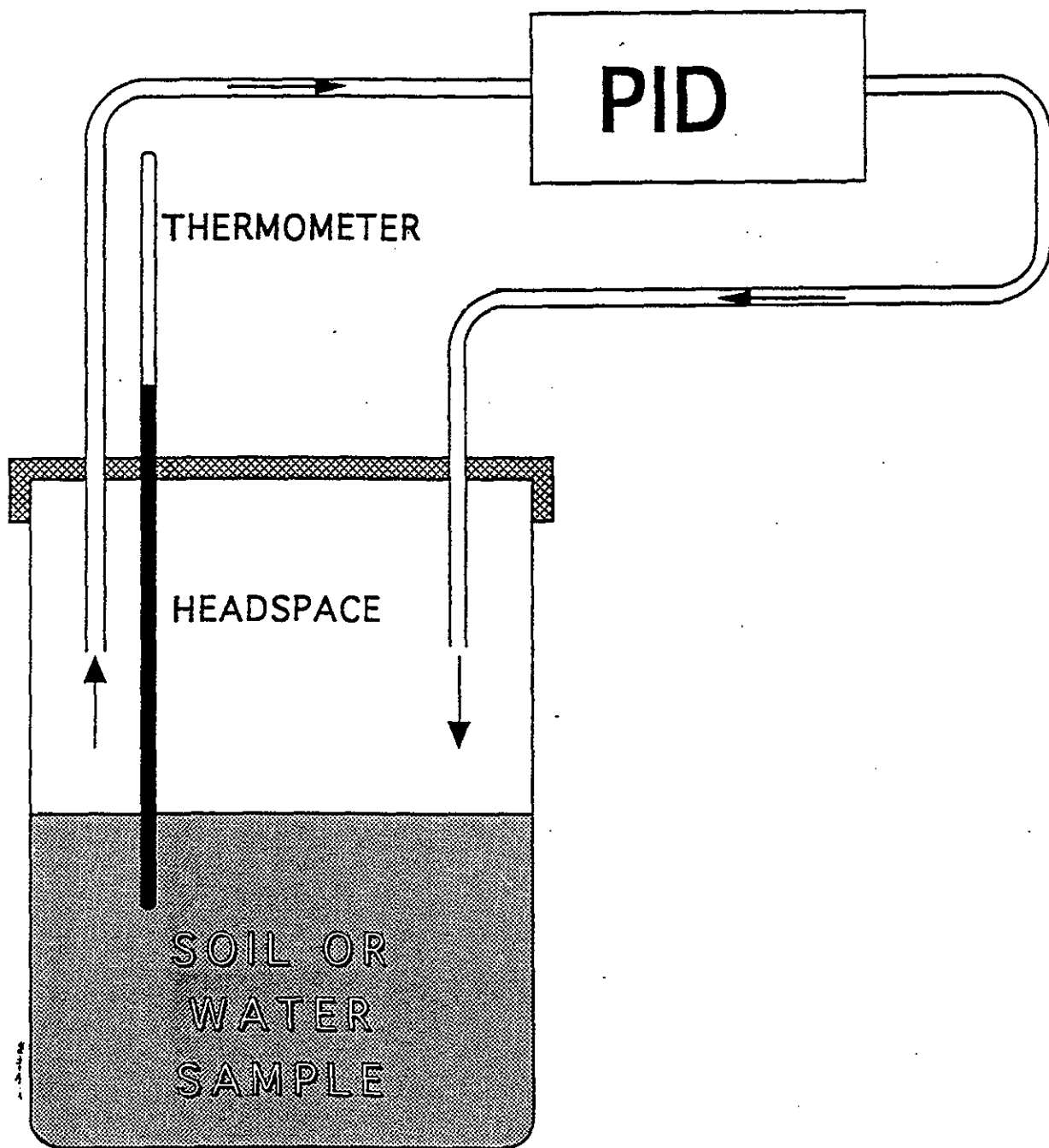
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**Table 1**  
**Equilibrium Headspace Measurements**  
**Well 699-S6-E4A**  
**September 27, 1996**

sample	collection time	analysis time	results
B0J281	0916	0930	3.2 ppm (OVM) / 17.2 ppm ( $\mu$ TIP)
B0J282	0936	1018	4.8 ppm ( $\mu$ TIP)
B0J283	0950	1030	5.3 ppm ( $\mu$ TIP)
B0J284	1005	Unable to obtain further readings due to failure of $\mu$ TIP (water drawn into detector cell) and lack of sufficient response from OVMs	
B0J285	1035		
B0J286	1105		
B0J287	1135		
B0J288	1205		



GENERAL CONCEPT OF THE EQUILIBRIUM  
HEADSPACE METHOD

## FIELD MONITORING DATA

Page \_\_\_\_\_ of \_\_\_\_\_

Instrument Type OVM / Colorimetric Tubes Identifier No. \_\_\_\_\_ Date 9/18/96  
 Project Name Well 699-S6-E4A Operators Name (Print/Sign) R.G. McCain

Well 699-S6-E4A

8.32 On site

Joe Jimenez, site super

7.18 completed pre-job safety &amp; RMP briefing

Kurt Higbee (TMA) is IH rep

OVM w 10.6 eV lamp calibrated to 97 ppm isobutylene

8.32 ready to uncap well

8.37 OVM under cap: no reading

8.38 uncap well OVM reads 0

8.39 ~ 5ft tygon tubing; OVM reads 1 ppm @ 5 ft

8.40 collecting Tedlar bag from tubing ~5 ft down well casing

8.41 OVM = 3.2-3.6

8.42 OVM = 4-4.9

8.43 OVM = 5.3

8.44 OVM = 5.3-5.7

8.45 OVM = 6.2

8.46 OVM = 5.7-6.2

8.47 OVM = 6.2-6.6

8.48 OVM = 6.2-6.6

8.49 OVM = 6.2-6.6

8.50 OVM = 6.2-6.6

8.51 OVM = 6.2-6.6

8.52 stop collecting bag

8.53 OVM = 6.6 @ ~3ft

8.54 OVM = 4 @ surface

8.55 OVM = 3.20

8.56 OVM bkgnd = 1.9-2.3

8.57 leave site; transport Tedlar bag to 100N in mobile lab

4

11.28 calibrate OVM (10.6 eV lamp) to 100 ppm isobutene

11.28 10 ppm isobutene = 7.8-10.9 ppm

11.30 bkgnd w needle on OVM is 0.5 ppm

11.33 10 ppm isobutene reads 6.8 ppm thru septum

11.35 unknown reads 0.5-1.5 ppm on OVM

11.42 Sensidyne Polytest (Exp9405): psble slight stain 1st 1-2 mm

11.52 Draeger Polytest; (Exp9412) slit ring stain 2mm to 4mm 1st brn

12.09 Sensidyne ethyl acetate (Exp7309): no stain

12.22 Draeger methyl bromide (Exp9410): no stain

12.31 Draeger polytest (Exp9412) w slit ring stain

12.33 connect OVM to bag: 0.5-1.5 ppm

12.38 OVM to 10 ppm isobutene in bag: 3.8 ppm

12.42 Draeger acetone (Exp9209) no stain

12.52 Sensidyne stoddard solvent (Exp9409) no stain

13.03 Sessle extended, unable to make any determination, probably because colorimetric tubes are too far out of date

## FIELD MONITORING DATA

Page \_\_\_\_\_ of \_\_\_\_\_

Instrument Type OVM, Equilibrium Headspace Identifier No. \_\_\_\_\_ Date 9/19/96  
 Project Name Well 699-S6-E4A Operators Name (Print/Sign) B.G. McLain

96-09-19

699-S6-E4A Well

On site @ 16.00

Dave St John & Doug Bryant are samplers  
 Kurt Higbee is SSG

Plan is to collect samples from the well with a bailer immediately  
 after perforation. Asked Dave to fill a 1-pint Mason jar ~ 1/2  
 full, in addition to other samples shown on GAF

set up OVM for equilibrium headspace measurements using 1-pint  
 Mason jars

16.39 Cal OVM to 100 ppm isobutene

16.41 10 ppm isobutene reads 9.6 ppm in headspace setup

17.20 began sample collection

dropped bailers approx 7 ft about 5 times  
 OVM reads 1.6 ppm @ top of casing  
 observed screen on water surface  
 water is a light dirty brown

17.46 received sample

water is light brown, cloudy, has small silt-size particles only  
 rust scale

17.48 start autolog mode on OVM 5 sec intervals

17.49 OVM reads 0.5 ppm on empty jar

17.51 OVM = 3.9 - 5.0 ppm on sample temp = 21 deg C

17.54 OVM reads 0.5 - 1.1 over DI water

17.56 OVM = 2.7 ppm over sample

18.07 turnoff autolog

18.07 add small qty Sudan IV dye to ~ 25 ml water. Shake &  
 centrifuge 2000 rpm for 5 min Dye is undissolved; no non aqueous  
 organic phase present

18.50 received sample 803202 2 500 ml amber wa glass + 1 500 ml  
 amber wa glass + HNO3 + 1 40 ml VGA vial

## FIELD MONITORING DATA

Page 1 of 3

Instrument Type \_\_\_\_\_

Identifier No. \_\_\_\_\_

Date 9/27/96Project Name Well 699-S6-E4AOperators Name  
(Print/Sign) B.G. McLain

96-09-27

699-S6-E4A

Well Development Activities

SAF: 896-187

6.30 on site

Doug Ewers &amp; Doug Bryant are samplers; Kurt Higbee is SSO

Procedure for field screening for VOCs (Equilibrium Headspace Method)

1.) Calibrate OVM to 100 ppm isobutene. Set average interval at 1 sec. Set auto-log on, with logging interval of 10 sec.

2.) Collect water in wat (475 ml) wide-mouth canning jar. Fill to depth of approximately 2.5 inches - volume of water is about 275 ml (42% headspace volume). Cap with retail canning lid.

3.) Replace the canning lid with the teflon headspace lid. Circulate headspace through the OVM and allow the reading to stabilize. Record the stable reading in ppm.

4.) After positive readings are obtained, remove the jar and allow the OM to recirculate ambient air until readings have stabilized at or near zero. Place a clean jar on the headspace lid and recirculate to ensure that no organic vapors remain in the headspace apparatus.

Equipment:

OVM1: K332501 ser# 5003-33358-243 10.6 eV lamp

OVM2: K335040 ser# 5000-35392-250 10.6 eV lamp

MicroTip HL2000 ser# 94920275

COI: GasTech "TraceTector" Ser#DT057

spar gas: 100 ppm isobutylene Byrne lot# 3-043 3/1/93

cal gas: 10 ppm isobutylene Byrne lot# 3-047 3/1/93

8.15 calibrate OVM1, OVM2, &amp; MicroTip to 100 ppm isobutylene

8.23 check response to 10 ppm isobutylene

OVM1 = 12.5 ppm OVM2 = 10.9-12.0

MicroTip = 10.1-10.2

8.45 check COI: 100 ppm isobutylene reads 79 ppm

9.15

9.17 Pump on

9.19 sample #1 200031 (samplers recorded time @ 9.18

9.20 Kurt Higbee reports no response on either OVM or GasTech

## FIELD MONITORING DATA

Page 2 of 3

Instrument Type \_\_\_\_\_ Identifier No. \_\_\_\_\_ Date 9/27/96  
 Project Name Well 699-56-E4A Operators Name (Print/Sign) B.G. McCain

9.21 flow reported at 6.1 gpm  
 9.29 Blank (DI water) reads 0.5 ppm (17C)  
 9.30 B0J281 reads 2.8-4.0 ppm (18C)  
 9.34 switch to OVM#1 reads 3.2 ppm (seems more stable) (19C)  
 9.39 MicroTip reads 17.2 ppm (19C)  
 9.42 rec'd 2nd sample from Doug Bryant B0J282, collected 9.31  
 9.50 OVM#1 reads 2.3 ppm on ambient air, MicroTip reads 1.1 ppm  
 10.07 rec'd 3rd sample from Doug Bryant B0J283  
 10.10 MicroTip reads 1.1 ppm on empty jar  
 10.12 MicroTip reads 2.0 ppm on DI water  
 10.14 rec'd 4th sample B0J283  
 10.18 B0J282 reads 4.8 ppm on microtip  
 10.25 add 31 ml B0J281 to centrifuge tube w pinch Gula IV, shake for 1 min & centrifuge @ 2000 rpm 10 min. Yields ~1 ml sediment with no evidence of a NAPL.  
 10.27 MicroTip reads 1.1 ppm on ambient air  
 10.30 B0J283 reads 5.5 ppm on MicroTip (21C)  
 10.32 pH of B0J281 is 9-10, conductivity is 412 uS/cm @ 19.5C  
 10.35 MicroTip reads 1.6 ppm.  
 10.43 fault on MicroTip: light intensity low  
 10.55 10 ppm isobutylene reads 13.2 ppm on OVM#2  
 10.56 unable to get a reading on B0J281 with OVM#2  
 Unable to get VOC readings with either OVM. MicroTip has drawn in a drop of water and is not functioning. Took MicroTip apart & dried out detector assembly, but unable to restore function. Still appears to be arcing inside detector housing.  
 LC Hulstrom has received preliminary U data from 1 st 3 samples. He has decided to terminate purging and collect samples for offsite analysis. Last screening sample collected at 12.05 Samples (583) delivered to Dr Jacques at 13.20 for Uranium analysis by KPA  
 Further examination of the water reveals the following



## FIELD MONITORING DATA

Page \_\_\_\_\_ of \_\_\_\_\_

Instrument Type \_\_\_\_\_ Identifier No. \_\_\_\_\_ Date 9/19/96Project Name Well 69956-E4A Operators Name (Print/Sign) R.G. McLain

Samples have a whitish material only slightly denser than water  
Later samples have an odor that resembles diesel  
The pH of these samples is ~10

2  
3  
4  
4